

PATENT APPLICATION

METHOD OF MAKING A LAYERED COMPOSITE ELECTRODE/ELECTROLYTE

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CROSS-REFERENCE TO RELATED APPLICATIONS

This application is a divisional of U.S. Application No. 09/626,022, filed July 27, 2000, titled SOLID STATE ELECTROCHEMICAL DEVICE ELECTRODE ADDITIVES, which claims priority from U.S. Provisional Application No. 60/146,767, filed July 31, 1999, titled
10 SURFACE ADDITIVES FOR ENHANCED ELECTRODE PERFORMANCE, the disclosures of which are herein incorporated by reference for all purposes.

BACKGROUND OF THE INVENTION

This invention was made with government support in the course of or under prime contract No. DE-ACO3-76SF00098 between the U.S. Department of Energy and the
15 University of California. The government has certain rights in this invention.

The present invention relates generally to the field of electrochemical devices, and more particularly solid state electrochemical devices composed of one or more electrodes in contact with a solid state electrolyte and/or membrane.

Solid state electrochemical devices are often implemented as cells including two
20 electrodes, the anode and the cathode, and a dense solid electrolyte/membrane which separates the electrodes. In many implementations, such as in fuel cells and oxygen and syn gas generators, the solid membrane is an electrolyte composed of a material capable of conducting ionic species, such as oxygen ions, sodium ions, or hydrogen ions, yet has a low electronic conductivity. In other implementations, such as gas separation devices, the solid
25 membrane is composed of a mixed ionic electronic conducting material ("MIEC"). In each case, the electrolyte/membrane must be gas-tight to the electrochemical reactants. In all of these devices a lower total internal resistance of the cell results in improved performance.

The preparation of solid state electrochemical cells is well known. For example, a typical solid oxide fuel cell is composed of a dense electrolyte membrane of a ceramic oxygen
30 ion conductor, a porous anode layer of a ceramic, a metal or, most commonly, a ceramic-metal composite ("cermet"), in contact with the electrolyte membrane on the fuel side of the cell, and a porous cathode layer of an ionically/electronically-conductive metal oxide on the oxidant side of the cell. Electricity is generated through the electrochemical reaction between a fuel (typically hydrogen produced from reformed methane) and an oxidant (typically air).

This net electrochemical reaction involves mass transfer and charge transfer steps that occur at the interface between the ionically-conductive electrolyte membrane, the electronically-conductive electrode and the vapor phase (fuel or oxygen). The contribution of these charge transfer steps, in particular the charge transfer occurring at the oxygen electrode, to the total internal resistance of a solid oxide fuel cell device can be significant.

Electrode structures including a porous layer of electrolyte particles on a dense electrolyte membrane with electrocatalyst material on and within the porous layer of electrolyte are known. As shown in Fig. 1, such electrodes are generally prepared by applying an electrocatalyst precursor-containing electrode material 102 (such as a metal oxide powder having high catalytic activity and high reactivity with the electrolyte) as a slurry to a porous (pre-fired; unsintered; also referred to as “green”) electrolyte structure 104, and then co-firing the electrode and electrolyte materials to densify the electrolyte and form a composite electrolyte/electrode/electrocatalyst 106.

Oxides containing transition metals such as Co, Fe, Mn, are known to be useful as oxygen electrodes in electrochemical devices such as fuel cells, sensors, and oxygen separation devices. However, if such compounds were to be used with typical zirconia-based electrolytes, such as YSZ, a deleterious reaction in the temperature range of 1000-1400°C typically needed to densify zirconia would be expected. The product of this reaction would be a resistive film 105 at the electrode/electrolyte interface, thereby increasing the cell’s internal resistance.

Similar problems may be encountered with sintering highly catalytic electrode materials on densified (fired) zirconia-base electrolytes since the sintering temperatures of about 1200 °C to 1400 °C are sufficient to cause the formation of a deleterious resistive film at the electrode/electrolyte interface.

In order to avoid deleterious chemical reactions, attempts have been made to use barrier layers, such as ceria, or to use chemically compatible electrolytes, such as ceria with such transition metal oxides. Also, it has been proposed to add an electrocatalytic precursor to a fired electrode/electrolyte composite, but only for a specific type of electrode material. Specifically, prior researchers have sought to fabricate electrodes with interpenetrating networks of ionically conductive and electronically conductive materials with subsequent infiltration of a catalytic electrode. However, this can hinder the performance of ionic devices, particularly at high current densities where ohmic drop due to current collection can lead to substantial efficiency losses.

It would be desirable to have improved techniques for fabricating high performance solid state electrochemical device oxygen electrodes composed of highly conductive materials.

SUMMARY OF THE INVENTION

The present invention addresses this need by providing an electrode fabricated from highly electronically conductive materials such as metals, metal alloys, or electronically conductive ceramics. In this way, the electronic conductivity of the electrode substrate is maximized. The electrode is combined with an electrolyte to form an electrode/electrolyte structure. The electrode/electrolyte structure of the invention may be prepared by a plurality of methods. An unsintered (possibly bisque fired) moderately catalytic electronically-conductive or homogeneous mixed ionic electronic conductive electrode material may be deposited on a layer composed of a sintered or unsintered ionically-conductive electrolyte material prior to being sintered. In one embodiment, a layer of particulate electrode material is deposited on an unsintered ("green") layer of electrolyte material and the electrode and electrolyte layers are sintered simultaneously, sometimes referred to as "co-firing," under conditions suitable to fully densify the electrolyte while the electrode retains porosity. In another embodiment, the layer of particulate electrode material is deposited on a previously sintered layer of electrolyte, and then sintered. Subsequently, a catalytic material is added to the electrode structure by infiltration of an electrocatalyst precursor (e.g., a metal salt such as a transition metal nitrate). This may be followed by low temperature firing to convert the precursor to catalyst. The invention allows for an electrode with high electronic conductivity and sufficient catalytic activity to achieve high power density in an ionic (electrochemical) device such as fuel cells and electrolytic gas separation systems.

In one aspect, the present invention provides a method of preparing a layered composite electrode/electrolyte structure. The method involves contacting a mixture of particles of an electronically-conductive or a homogeneous mixed ionically electronically-conductive (MIEC) electrode material with a layer of an ionically-conductive electrolyte material to form an assembly having a layer of the mixture on at least one side of the layer of the electrolyte material. The assembly is sintered to form a porous electrode in contact with a gas-tight electrolyte membrane. The sintered assembly is then infiltrated with a solution or dispersion of a transition metal nitrate-containing electrocatalyst precursor.

In another aspect, the invention provides a method of preparing a layered composite electrode/electrolyte structure. The method involves contacting a mixture of particles of a metal alloy selected from the group consisting of a chromium-containing ferritic steel, a chrome-based alloy and a chrome-containing nickel-based alloy with a layer of an ionically-conductive electrolyte material to form an assembly having a layer of the mixture on at least one side of the layer of the electrolyte material. The assembly is sintered to form a porous electrode in contact with a gas-tight electrolyte membrane. The sintered assembly is then infiltrated with a solution or dispersion of an electrocatalyst precursor.

These and other features and advantages of the present invention will be presented in more detail in the following specification of the invention and the accompanying figures which illustrate by way of example the principles of the invention.

BRIEF DESCRIPTION OF THE DRAWINGS

The present invention will be readily understood by the following detailed description in conjunction with the accompanying drawing:

Fig. 1 depicts an undesirable method of forming an electrode/electrolyte composite for a solid state electrochemical cell.

Fig. 2 depicts stages in a method of forming an electrode/electrolyte composite for a solid state electrochemical cell in accordance with one embodiment the present invention.

Fig. 3 shows plots of the impedance spectra for an electrode with and without additives in accordance with one embodiment of the present invention.

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DETAILED DESCRIPTION OF THE INVENTION

Reference will now be made in detail to some specific embodiments of the invention including the best modes contemplated by the inventors for carrying out the invention. Examples of these specific embodiments are illustrated in the accompanying drawings.

5 While the invention is described in conjunction with these specific embodiments, it will be understood that it is not intended to limit the invention to the described embodiments. On the contrary, it is intended to cover alternatives, modifications, and equivalents as may be included within the spirit and scope of the invention as defined by the appended claims. In the following description, numerous specific details are set forth in order to provide a
10 thorough understanding of the present invention. The present invention may be practiced without some or all of these specific details. In other instances, well known process operations have not been described in detail in order not to unnecessarily obscure the present invention.

15 In general, the present invention provides a technique for the addition of highly reactive elements known to be of value for enhancing catalytic properties of the oxygen electrode to the surface where they are needed, without chemical reactions that occur during firing on of these electrode materials. In this way, moderate performance cathodes that are known to be more chemically stable at the firing temperatures can be used for the electrode's microstructure formation. Then high performance additives can be incorporated into the
20 electrodes without significantly altering the microstructure or creating performance limiting reaction layers.

In this specification and the appended claims, the singular forms "a," "an," and "the" include plural reference unless the context clearly dictates otherwise. Unless defined otherwise, all technical and scientific terms used herein have the same meaning as commonly
25 understood to one of ordinary skill in the art to which this invention belongs.

For the purposes of this application, an oxygen (or air) electrode refers to the electrode at which oxygen is either reduced, or oxygen anions are oxidized, depending on the function of the cell. Examples of oxygen electrodes are the cathode portion of a solid oxide fuel cell or the anode portion of an electrolytic cell, such as may be used for oxygen generation. The
30 oxygen electrode portion of the electrode/electrolyte structure of this invention comprises a porous, electronically-conductive material, or a (homogeneous) mixed ionic electronic conductor ceramic material having an electrocatalyst dispersed within its pores.

Fig. 2 show stages in a method of forming an electrode/electrolyte composite for a solid state electrochemical cell in accordance with one embodiment the present invention.

Referring to Fig. 2(A), a particulate electrode material 202 is applied as a slurry to a porous green (pre-fired; unsintered) electrolyte structure 204 to form an assembly 206.

The electrode/electrolyte structure of the invention may be prepared by any suitable method. For example, an unsintered (possibly bisque fired) moderately catalytic electronically-conductive or homogeneous mixed ionic electronic conductive electrode material may be deposited on a layer composed of a sintered or unsintered ionically-conductive electrolyte material prior to being sintered. In one embodiment, a layer of particulate electrode material is deposited on an unsintered layer of electrolyte material and the electrode and electrolyte layers are sintered simultaneously, sometimes referred to as “co-firing.” In another embodiment, the layer of particulate electrode material is deposited on a previously sintered layer of electrolyte, and then sintered.

The particulate electrode material may be applied to the electrolyte layer (generally composed of particles of an electrolyte material), by any suitable means such as, for example, aerosol spray, dip coating, painting or silk-screening, electrophoretic deposition, vapor deposition, vacuum infiltration, and tape casting.

In general, suitable electrode materials in accordance with the present invention as have an electronic conductivity of at least about 10^{-1} Siemens/cm (S/cm) at the operating temperature of the electrochemical device with which the electrode is to be incorporated. Preferably, the conductivity of the material is at least about 100 S/cm, more preferably at least about 1000 S/cm. The electrode material should also be compatible with the electrolyte layer; that is, it should not cause any deleterious reaction with the electrolyte at the processing temperatures to form a new phase with insufficient conductivity or electrocatalytic properties for use in a practical electrochemical device.

Exemplary electronically-conductive materials include metals, such as the transition metals Cr, Fe, Cu, Ag, and metal alloys thereof. Such metal electrode materials are suitable for use in oxygen or hydrogen electrodes for solid state electrochemical devices that are operated at relatively low temperatures, for example, about 400 to 800°C.

Other suitable electrode materials include mixed ionic electronic conductor ceramic materials, such as $\text{La}_{1-x}\text{Sr}_x\text{Mn}_y\text{O}_{3-\delta}$ ($1 \geq x \geq 0.05$) ($0.95 \leq y \leq 1.15$) (“LSM”), $\text{La}_{1-x}\text{Sr}_x\text{CoO}_{3-\delta}$ ($1 \geq x \geq 0.10$) (“LSC”), $\text{SrCo}_{1-x}\text{Fe}_x\text{O}_{3-\delta}$ ($0.30 \geq x \geq 0.20$), $\text{La}_{0.6}\text{Sr}_{0.4}\text{Co}_{0.6}\text{Fe}_{0.4}\text{O}_{3-\delta}$, $\text{Sr}_{0.7}\text{Ce}_{0.3}\text{MnO}_{3-\delta}$, $\text{LaNi}_{0.6}\text{Fe}_{0.4}\text{O}_3$, and $\text{Sm}_{0.5}\text{Sr}_{0.5}\text{CoO}_3$. Preferred LSM materials include $\text{La}_{0.8}\text{Sr}_{0.2}\text{MnO}_{3-\delta}$, $\text{La}_{0.65}\text{Sr}_{0.35}\text{MnO}_{3-\delta}$ and $\text{La}_{0.45}\text{Sr}_{0.55}\text{MnO}_{3-\delta}$. These materials form a homogeneous electrode having both ionic and electronic conductivity.

Low-chromium ferritic steels, such as type 405 and 409 (11-15% Cr), intermediate-chromium ferritic steels, such as type 430 and 434, (16-18% Cr), or high-chromium ferritic steels, such as type 442, 446 and E-Brite (19-30% Cr), chrome based alloys such as Cr5Fe1Y, and chrome-containing nickel-based Inconel alloys including Inconel 600 may also be used.

5 Suitable electrolyte/membrane materials for use in solid state electrochemical devices are well known in the art. In general, they are ionically-conductive solid membranes having an ionic conductivity of at least about 10^{-3} S/cm at the operating temperature of the device and sufficiently low electronic conductivity for use as the electrolyte membrane which separates the anode from the cathode in a solid state electrochemical device. Doped zirconias
10 such as yttria-stabilized zirconia ("YSZ") and scandium-doped zirconia are very commonly used. Preferably, the electrolyte/membrane material is YSZ.

In the case of an electrolytic oxygen separation device, where oxygen is driven across the membrane by applying a potential difference and supplying energy, the membrane may be chosen from electrolytes well known in the art including yttria stabilized zirconia (e.g.,
15 $(\text{ZrO}_2)_{0.92}(\text{Y}_2\text{O}_3)_{0.08}$), scandia stabilized zirconia (SSZ), doped ceria such as $(\text{CeO}_2)_{0.8}(\text{Gd}_2\text{O}_3)_{0.2}$ (CGO), doped lanthanum gallate such as $\text{La}_{0.8}\text{Sr}_{0.2}\text{Ga}_{0.85}\text{Mg}_{0.15}\text{O}_{2.825}$ (LSGM20-15) and doped bismuth oxide such as $(\text{Bi}_2\text{O}_3)_{0.75}(\text{Y}_2\text{O}_3)_{0.25}$.

In the case of a gas separation device, where partial pressures, rather than applied potential, are used to move ions across the membrane, the membrane may be a mixed ionic
20 electronic conductor (MIEC), such as $\text{La}_{1-x}\text{Sr}_x\text{CoO}_{3-\delta}$ ($1 \geq x \geq 0.10$) ("LSC"), $\text{SrCo}_{1-x}\text{Fe}_x\text{O}_{3-\delta}$ ($0.30 \geq x \geq 0.20$), $\text{La}_{0.6}\text{Sr}_{0.4}\text{Co}_{0.6}\text{Fe}_{0.4}\text{O}_{3-\delta}$, $\text{Sr}_{0.7}\text{Ce}_{0.3}\text{MnO}_{3-\delta}$, $\text{LaNi}_{0.6}\text{Fe}_{0.4}\text{O}_3$, and $\text{Sm}_{0.5}\text{Sr}_{0.5}\text{CoO}_3$.

Such electrolyte/membranes are well known to those of skill in the art and may be prepared by any suitable method, such as by depositing a slurry of an ionically-conductive electrolyte material directly onto one of the electrodes, or by preparing a cast tape of an
25 ionically-conductive electrolyte material, which is laminated to a cast tape of electrode material.

The assembly 206 is then fired at a temperature and under conditions suitable to sinter the electrode material, densify the electrolyte (where necessary), and bond the two to form a composite electrolyte/electrode 208, as shown in Fig. 2(B). The sintering conditions should
30 be selected so that they are sufficient to fuse the majority of the electrode particles. If the electrolyte is green, the sintering conditions should be selected to densify the electrolyte material sufficiently to form a gas-tight electrolyte membrane. Suitable sintering conditions for given materials may be readily determined experimentally by those of skill in the art. When sintered, the electrode material forms a porous layer.

As shown in Fig. 2(C), an electrocatalytic precursor is then incorporated into the porous sintered electrode 210. The electrocatalytic precursor is added by any suitable technique, such as by infiltrating the network with a solution or dispersion of an electrocatalyst precursor and heating the infiltrated network under conditions sufficient to form the corresponding electrocatalyst. The electrocatalyst should have sufficient catalytic activity for the electrochemical reaction(s) occurring at the electrode for its use in a practical device. This material must also be compatible with the electrolyte layer at the operating temperature of the device.

Suitable electrocatalytic precursors in accordance with the present invention include aqueous or non-aqueous solutions of metal salts, such as nitrates, acetates and citrates. For example, this can be accomplished by using nitrate salts of a transition or rare earth metal or combination of transition and/or rare earth metal salts desired for catalytic activity, such as Co nitrate, Fe nitrate, Mn nitrate or, for a hydrogen electrode, Ni nitrate. The porous electrode structure may be infiltrated by any suitable means such as by aerosol spray, dip coating, painting or silk-screening, or vacuum infiltration of the electrocatalyst material into the porous structure. A stack of cells may also be assembled prior to being infiltrated and infiltrated simultaneously.

Following addition of the electrocatalytic precursor, the composite electrode/electrolyte 212 is heated to operating temperature for the electrochemical device of which it is a part, generally about 600 to 900°C, as shown in Fig. 2(D). In this way a highly catalytic electrode in a composite electrode/electrolyte structure may be produced without deleterious reactions between the electrocatalytic precursors and the zirconia of the electrolyte at the electrode/electrolyte bonding production stage. Thus, in one example, an LSM/YSZ structure can be fired at reasonably high temperatures (1350°C) to yield a moderate performance electrode, and by adding cobalt nitrate by infiltration to this electrode the performance is boosted to what might be expected for cobalt based electrode.

The porosity of the composite electrode structure containing the electrocatalyst is preferably at least about 10 percent, more preferably at least about 20 percent, and generally about 30%. The thickness of the sintered electrolyte membrane is preferably between about 5 and 40 microns, more preferably no greater than about 20 microns. The thickness of the sintered composite electrode is preferably between about 5 and 5000 microns, more preferably no greater than about 2000 microns (a thicker electrode is used where the electrode also provides the mechanical support for an electrochemical device).

Infiltration of a fired electrode with a electrocatalytic precursor solution (e.g., a metal nitrate solution) not only solves the problem of reactivity of the catalytic material with the electrolyte, but also addresses the problem of thermal mismatch. If, for example, a highly

catalytic electrode material had a very poor match of the coefficient of thermal expansion with the electrolyte, thermal cycling would typically lead to spallation of the electrode from the electrolyte surface. In the case of the present invention, an electrode material that has a thermal expansion coefficient close to that of the electrolyte may be used. The electrode is therefore well bonded to the electrolyte. In a subsequent step, an electrocatalytic precursor for a highly catalytic material, such as such as a metal oxide, is infiltrated into the bonded electrode thereby improving the electrode performance.

Importantly, this process represents a simple low-cost means of improving electrode performance by allowing the fabrication of a high strength, high conductivity electrode from materials of moderate catalytic activity and then adding that activity in a subsequent step. Furthermore, in the case of expensive catalytic materials, a low cost material can be used to develop the electrode microstructure followed by a small amount of catalytic material introduced by the metal salt technique described herein.

Example

The following example illustrates aspects and features of a specific implementation in accordance with the present invention. It should be understood the following is representative only, and that the invention is not limited by the detail set forth in this example.

A 1cm x 1cm Pt electrode (previously fired to 950°C to bond Pt to electrolyte) was tested on YSZ electrolyte at 900°C. LSM-YSZ was used as the counter electrode. Then a mixture of Co, Sr, Ce, and Ni nitrates dissolved in distilled water were applied to the Pt electrode and allowed to dry under a heat lamp. The sample was again placed in a furnace and heated. Impedance spectra was taken at 900°C. Fig. 3 shows plots of the impedance spectra for the Pt electrode without additives (large arc) and Pt electrode with surface additives (small arc). The example shows the effect of a bare porous metal electrode, Platinum, with and without the addition of metal nitrate salts. The reduction in the size of the impedance arc for the electrode with the additives shows improvement in performance by lowering cell impedance.

CONCLUSION

Although the foregoing invention has been described in some detail for purposes of clarity of understanding, those skilled in the art will appreciate that various adaptations and modifications of the just-described preferred embodiments can be configured without departing from the scope and spirit of the invention. Moreover, the described processing distribution and classification engine features of the present invention may be implemented

together or independently. Therefore, the described embodiments should be taken as illustrative and not restrictive, and the invention should not be limited to the details given herein but should be defined by the following claims and their full scope of equivalents.

What is claimed is: